

AMEL - Observational study of the management of metastatic melanoma in France

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Last update : 10/19/2017 | Version : 1 | ID : 73340

General	
Identification	
Detailed name	Observational study of the management of metastatic melanoma in France
Sign or acronym	AMEL
General Aspects	
Medical area	Cancer research
Pathology (details)	metastatic melanoma
Scientific investigator(s) (Contact)	
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Unit	MSD France
Collaborations	
Funding	
Funding status	Private
Details	MSD France
Governance of the database	
Sponsor(s) or organisation(s) responsible	MSD France
Organisation status	Private
Additional contact	

Main features

Type of database

Type of database

Study databases

Study databases (details)

Longitudinal study (except cohorts)

Database objective

Main objective

Describe the sociodemographic and clinical profile of patients with metastatic melanoma as well as their therapeutic management

Inclusion criteria

Inclusion criteria:

? History of histopathologically confirmed diagnosis of cutaneous melanoma

? Unresectable stage III (per treating physician) or stage IV melanoma according to AJCC (2009) classification at diagnosis or upon progression following diagnosis

? Subject's index date is between 1 January 2012 and 31 October 2012.

? Age \geq 18 years at the index date

? Access to the complete records from the index date until the end of the data extraction period or death whichever is earlier

Non-inclusion criteria:

? Patients with primary non-cutaneous melanoma including primary ocular melanoma

? Patients with an index date before 1 January 2012 and 31 October 2012.

? Patients with resectable melanoma who have not relapsed

Population type

Age

Adulthood (19 to 24 years)

Adulthood (25 to 44 years)

Adulthood (45 to 64 years)

Elderly (65 to 79 years)

Great age (80 years and more)

Population covered

Sick population

Pathology

C00-C97 - Malignant neoplasms

Gender

Male

Woman

Geography area

National

Detail of the geography area	no information about the individual
Data collection	
Dates	
Size of the database	
Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	271 patients included
Data	
Database activity	Data collection completed
Type of data collected	Clinical data Administrative data Cost data
Clinical data (detail)	Direct physical measures
Presence of a biobank	No
Procedures	
Participant monitoring	No
Followed pathology	
Links to administrative sources	No
Promotion and access	
Promotion	
Access	
Terms of data access (charter for data provision, format of data, availability delay)	No data access charter
Access to aggregated data	Access not yet planned
Access to individual data	No access